PTO/SB/06a (05-07)
Approved for use through 09/30/2007 CMB 0651-0031
U.S. Patent and Trademark Office, U.S. DEPARTIMENT OF COMMERCE to a collection of information unless it contains a valid OMS control number. Under the Paperwork Reduction Act of 1995, no persons are requir

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
( Not for submission under 37 CFR 1.99)

Application Number		10540384
Filing Date		2006-01-04
First Named Inventor Sum		Naoki
Art Unit		2823
Examiner Name	Maide	onado, Julio J.
Attorney Docket Number		1176/305

				Attorn	ey Doc	ket Number		1176/305				
					U.S.	PATENTS				Remove		_
Examiner Initial*	Cite No	Patent Number	Kind Code <sup>1</sup>	Issue E	Issue Date Name of Patentee or Applicant		Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear					
	1											
If you wisl	h to a	l dd additional U.S. Pater	nt citatio	n inform	ation pl	lease click the	Ac	id button.		Add		_
			U.S.P	ATENT	APPLI	CATION PUB	LIC	CATIONS		Remove		
Examiner Initial*	Cite No	Publication Number	Kind Code <sup>1</sup>	Publica Date	Name of Patentee or Applicant of cited Document			Relev	s,Columns,Line ant Passages as Appear			
	1	20020196517		2002-12	1-26	Nimura						
If you wisl	h to a	dd additional U.S. Publi	shed Ap	plication	citatio	n information p	ole	ase click the Ad	butto	n. Add		
				FOREIG	SN PAT	TENT DOCUM	E	NTS		Remove		
Examiner Initial*	Cite No	Foreign Document Number <sup>3</sup>	Country Code <sup>2</sup>		Kind Code <sup>4</sup>	Publication Date	A	lame of Patente opplicant of cited occument		Pages,Colum where Releva Passages or F Figures Appe	nt Relevant	TE
	1	4199681	JP			1992-07-20	F	ujitsu Lld.				
	2	1381825	CN			2002-11-27		oujing Science & echnology				
If you wisi	h to a	dd additional Foreign P	atent Do	cument	citation	I information pl	l lea	se click the Add	buttor	Add		
			NON	I-PATE	NT LITE	RATURE DO	CL	JMENTS		Remove		_

#### 

Examiner Initials*	Cite No	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, city and/orc country where published.	Ţ5
	1		

If you wish to add additional non-patent literature document citation information please click the Add button Add

# Examiner Signature Date Considered

\*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

1 See Kind Codes of USPTIO Patent Documents at year LISPTIO, CODE or MEPE 901.6.2 Elector office that issued the document, by the holietier code (WIPO Standard ST.3). 3 For disparsee patent obcurreds, by the indication of the year of the Propriet many percent the section number of the Propriet many percent the section number of the Propriet Standard ST.16 if possible. 3 Applicant is to place a check mark here if Explaint Indication should be controlled to the document under WIPO Standard ST.16 if possible. 3 Applicant is to place a check mark here if

## INFORMATION DISCLOSURE STATEMENT BY APPLICANT ( Not for submission under 37 CFR 1.99)

Application Number		10540384		
Filing Date		2006-01-04		
First Named Inventor Sumi,		Naoki		
Art Unit		2823		
Examiner Name	Maide	nado, Julio J.		
Attorney Docket Number		1176/305		

#### CERTIFICATION STATEMENT

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s):

	That each item of information contained in the information disclosure statement was first cited in any communication
$\times$	from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the
	information displacate statement. Con 27 CED 1 07(a)/1)

OR

That no item of information contained in the information disclosure statement was cited in a communication from a
foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification
after making reasonable inquiry, no item of information contained in the information disclosure statement was known to
any individual designated in 37 CFR 1.56(c) more than three months prior to the filing of the information disclosure
statement. See 37 CFR 1.97(e)(2).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- ....

#### SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/Wen Liu, Reg. #32,822/	Date (YYYY-MM-DD)	2007-08-09
Name/Print	Wentin	Registration Number	32 822

This collection of information is required by 3T CFR 1.87 and 1.98. The information is required to obtain or retain a benefit by the public which is to file fand by the USPTO to process) an application. Confidentially is governed by \$5 U.S. C. 12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Tradenary (Tifice, U.S. perpartment of Commence, P. 0. Box 1445), Alexandria, V.S. 2231-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VAZ 2331-1450.

### Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the stacked form related to a petient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) familishing of the information solicided is civulating; and (3) the principal purpuse for which the information is used by the U.S. Patient and Trademan Coffice is to process and/or cosmisting your submission related to a patient agricultant or patient. If you do not furnish the requested process and/or cosmisting your submission related to a patient agricultant or patient. If you do not furnish the requested process and the process of the process and the process of the pro

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these cords.
  - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiations.
  - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record perfains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
  - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552(m).
  - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
    may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
    to the Patent Cooperation Treaty.
  - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
  - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an insection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 4d U.S.C. 2904 and 2905. Such disclosure shall be made in accordance with the GSA requisions governing inseption of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the
  application pursuant to 35 U.S.C. 12(2) to rissuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be
  disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filled in application
  which became abandoned or in which the proceedings were terminated and which application is referenced by either a
  published application, an application open to public inspections or as issued patent.
  - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.